

**Abstract of the Disclosure**

The present invention relates to an oligonucleotide conjugate, comprising: (a) an oligonucleotide at least part of whose sequence is complementary to an intracellular nucleic acid sequence; and (b) a somatostatin analog. The present invention also relates to a medicament containing this oligonucleotide conjugate, preferably for treating tumors in which the somatostatin receptor (SSTR) is overexpressed.